SNV #	SNV name	Chr.	Position (build 36)	Gene	Annotation	Alleles (effect/ other)	EAF	Unconditional		Conditional on SNV #1		Conditional on SNVs #1 and #2		LD with SNV #1	LD with SNV #2
								Effect	P	Effect	P	Effect	P	r ²	r ²
CUBN	/ region														
1	rs1801222	10	17,196,157	CUBN	F253S	G/A	0.593	0.11	2.3×10^{-42}						
2	rs56077122	10	17,247,021	TRDMT1	intronic	A/C	0.335	0.061		0.0866	4.8×10^{-21}			0.033	
TCN1	region														
1	rs34324219	11	59,379,954	TCN1	D301Y	C/A	0.889	0.21	9.8×10^{-62}						
2	rs34528912	11	59,388,111	TCN1	R35H	T/C	0.0361	0.15	2.5×10^{-12}	0.17	2.1×10^{-15}			0.0040	
3	rs117456053	11	59,373,407	Near TCN1	Intergenic	G/A	0.976	0.12	4.1×10^{-6}	0.15	1.2×10^{-8}	0.16	1.9 × 10 ⁻⁹	0.0035	0.0011
TCN2	region														
1	rs1131603	22	29,348,975	TCN2	L376S	C/T	0.055	0.17	1.1×10^{-21}						
2	rs5753231	22	29,333,069	TCN2	Promoter	C/T	0.79	0.053	1.9×10^{-7}	0.064	7.5×10^{-10}			0.014	
мтн	FR region														
1	rs1801133	1	11,778,965	MTHFR	A222V	G/A	0.668	0.10	3.4×10^{-27}						
2	rs17421511	1	11,780,375	MTHFR	Intronic	G/A	0.827	0.045	0.00011	0.098	1.8×10^{-15}			0.11	

Conditional analyses were performed using imputed sequence data from chip-typed Icelanders with information on serum B_{12} or folate levels. Results for SNV #1 (lead SNVs) at each loci are unconditional on other SNVs. Analysis of SNV #2 is conditional on SNV #1 and SNV #3 is conditional on SNV #1 and #2. The LD between the SNVs at each locus was estimated from the sequence information of the 1,179 whole genome sequenced Icelanders.